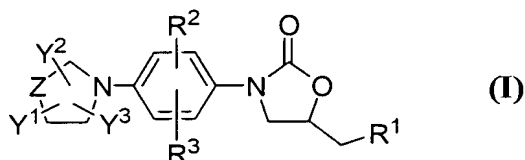


IN THE CLAIMS

1. (Previously Presented) A compound of the formula (I)



wherein

R^1 represents halo, azido, isothiocyanate, thioalcohol, OR^4 , NHR^4 or $N(R^4)_2$, where R^4 represents substituted or unsubstituted groups selected from acyl, thioacyl, (C_1-C_6) alkoxycarbonyl, (C_3-C_6) ycloalkoxythiocarbonyl, (C_2-C_6) alkenyloxycarbonyl, (C_2-C_6) alkenylcarbonyl, aryloxycarbonyl, (C_1-C_6) alkoxythiocarbonyl, (C_2-C_6) alkenyloxythio-carbonyl, aryloxythiocarbonyl, $-C(=O)-C(=O)-alkyl$, $-C(=O)-C(=O)-aryl$, $-C(=O)-C(=O)-alkoxy$, $-C(=O)-C(=O)-aryloxy$, $-(C=S)-S-alkyl$, $-(C=S)-NH_2$, $-(C=S)-NH-alkyl$, $-C(=S)-N-(alkyl)_2$, $-C(=S)-NH-alkenyl$, $(C=S)-(C=O)-alkoxy$, $-(C=S)-(C=O)-aryloxy$, $-C(=S)-O-(C=O)-alkyl$, $C(=S)-C(=S)-alkyl$, $-C(=S)-C(=S)-aryl$, thiomorpholinylthiocarbonyl or pyrrolidinylthiocarbonyl;

R^2 and R^3 are same or different and independently represent hydrogen, halogen atom, (C_1-C_6) alkyl group, halo (C_1-C_6) alkyl, cyano, nitro, SR^a , NR^a , OR^a where R^a represents substituted or unsubstituted (C_1-C_6) alkyl group, or halo (C_1-C_6) alkyl;

Z represents O,

Y^1 represents $=O$ or $=S$ group and Y^2 and Y^3 independently represent hydrogen, halogen, cyano, nitro, formyl, hydroxy, amino, $=O$, $=S$ group, or substituted or unsubstituted groups selected from (C_1-C_6) alkyl, hydroxy (C_1-C_6) alkyl, (C_1-C_6) alkoxy (C_1-C_6) alkyl, (C_1-C_6) alkoxycarbonyl, carboxy (C_1-C_6) alkyl, (C_1-C_6) alkylsulfonyl, (C_1-C_6) alkylcarbonyl-amino (C_1-C_6) lkyl, arylcarbonylamino (C_1-C_6) alkyl, (C_1-C_6) alkylcarbonyloxy (C_1-C_6) alkyl, amino (C_1-C_6) alkyl, mono (C_1-C_6) alkylamino, di (C_1-C_6) alkylamino, arylamino, (C_1-C_6) alkoxy, aryl, aryloxy, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocycloalkyl;

Y^2 and Y^3 when present on adjacent carbon atoms together also form a substituted or unsubstituted 5 or 6 membered aromatic or non-aromatic

cyclic structure, optionally containing one or two hetero atoms; its stereoisomers, its polymorphs, its pharmaceutically acceptable salts or its pharmaceutically acceptable solvates.

2. (Original) The compound according to claim 1, wherein the substituents on R⁴ are selected from halogen, hydroxy, amino, monoalkylamino, dialkylamino, cyano, nitro, alkoxy, aryl, hydroxyaryl, pyridyl, hydroxyalkyl, alkoxyaryl or carboxyl and its derivatives.

3. (Cancelled)

4. (Original) The compound according to claim 1, wherein the substituents on Y² and Y³ are selected from hydroxy, nitro, cyano, amino, *tert*-butyldimethylsilyloxy (TBSO), halogen, (C₁-C₆)alkyl, (C₁-C₆)alkoxy, (C₃-C₆)cycloalkyl, aryl, benzyloxy, acyl, carboxyl or acyloxy groups.

5. (Original) The compound according to claim 1, wherein the cyclic structure formed by Y² and Y³ is selected from substituted or unsubstituted benzene, pyridine, pyrrolidine, furan, thiophene, morpholine, piperazine or pyrrole.

6. (Previously Presented) A compound of the formula (I) as defined according to claim 1, which is selected from:

(5R)-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-5-hydroxymethyl-1,3-oxazolan-2-one or its salts;

(5R)-3-[3-fluoro-4-(2-thioxo-1,3-oxazolan-3-yl)phenyl]-5-hydroxymethyl-1,3-oxazolan-2-one or its salts;

3-{2-fluoro-4-[(5R)-5-hydroxymethyl-2-oxo-1,3-oxazolan-3-yl]phenyl}-2,3-dihydrobenzo[d][1,3]oxazol-2-one or its salts;

3-{2-fluoro-4-[(5R)-5-hydroxymethyl-2-oxo-1,3-oxazolan-3-yl]phenyl}-6-methyl-2,3-dihydrobenzo[d][1,3]oxazol-2-one or its salts;

3-{2-fluoro-4-[(5R)-5-hydroxymethyl-2-oxo-1,3-oxazolan-3-yl]phenyl}-5-methyl-2,3-dihydrobenzo[d][1,3]oxazol-2-one or its salts;

(5R)-5-hydroxymethyl-3-[4-(2-oxo-1,3-oxazolan-3-yl)-3-trifluoromethylphenyl]-1,3-oxazolan-2-one or its salts;

(5R)-3-[2-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-5-

hydroxymethyl-1,3-oxazolan-2-one or its salts;
(5R)-3-[3,5-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-5-hydroxymethyl-1,3-oxazolan-2-one or its salts;
(5R)-5-hydroxymethyl-3-[4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-1,3-oxazolan-2-one or its salts;
3-{4-[(5R)-5-hydroxymethyl-2-oxo-1,3-oxazolan-3-yl]phenyl}-2,3-dihydrobenzo[d][1,3]oxazol-2-one or its salts;
(5R)-azidomethyl-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-1,3-oxazolan-2-one or its salts;
3-{4-[(5R)-5-azidomethyl-2-oxo-1,3-oxazolan-3-yl]-2-fluorophenyl}-6-methyl-2,3-dihydrobenzo[d][1,3]oxazol-2-one or its salts;
3-{4-[(5R)-5-azidomethyl-2-oxo-1,3-oxazolan-3-yl]-2-fluorophenyl}-5-methyl-2,3-dihydrobenzo[d][1,3]oxazol-2-one or its salts;
(5R)-5-azidomethyl-3-[4-(2-oxo-1,3-oxazolan-3-yl)-3-trifluoromethylphenyl]-1,3-oxazolan-2-one or its salts;
(5R)-5-azidomethyl-3-[2-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-1,3-oxazolan-2-one or its salts;
(5R)-azidomethyl-3-[3,5-difluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-1,3-oxazolan-2-one or its salts;
(5R)-5-azidomethyl-3-[4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-1,3-oxazolan-2-one or its salts;
3-{4-[(5R)-5-azidomethyl-2-oxo-1,3-oxazolan-3-yl]phenyl}-2,3-dihydrobenzo[d][1,3]oxazol-2-one or its salts;
(5R)-aminomethyl-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-1,3-oxazolan-2-one or its salts;
(5R)-5-aminomethyl-3-[4-(2-oxo-1,3-oxazolan-3-yl)-3-trifluoromethylphenyl]-1,3-oxazolan-2-one or its salts;
(5R)-5-aminomethyl-3-[4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-1,3-oxazolan-2-one or its salts;
N-{(5S)-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} methanamide or its salts;
N-{(5S)-3-[3,5-difluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} methanamide or its salts;

N1-((5S)-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} acetamide or its salts;

N1-((5S)-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} propanamide or its salts;

N1-((5S)-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} butanamide or its salts;

N1-((5S)-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} pentanamide or its salts;

N1-((5S)-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} heptanamide or its salts;

N1-((5S)-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} acrylamide or its salts;

N1-((5S)-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl}-2,2,2-trifluoroacetamide or its salts;

Ethyl(5S)-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl carbamoylmethanoate or its salts;

N1-((5S)-3-[3-fluoro-4-(2-thioxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} acetamide or its salts;

N1-((5S)-2-oxo-3-[4-(2-thioxo-1,3-oxazolan-3-yl)phenyl]-1,3-oxazolan-5-ylmethyl} acetamide or its salts;

N1-((5S)-3-[3-fluoro-4-(2-oxo-2,3-dihydrobenzo[d] [1,3] oxazol-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} acetamide or its salts;

N1-((5S)-3-[3-fluoro-4-(6-methyl-2-oxo-2,3-dihydrobenzo[d] [1,3] oxazol-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} acetamide or its salts;

N1-((5S)-3-[3-fluoro-4-(5-methyl-2-oxo-2,3-dihydrobenzo[d] [1,3] oxazol-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} acetamide or its salts;

N1-((5S)-2-oxo-3-[4-(2-oxo-1,3-oxazolan-3-yl)-3-trifluoromethylphenyl]-1,3-oxazolan-5-ylmethyl} acetamide or its salts;

N1-((5S)-2-oxo-3-[4-(2-oxo-1,3-oxazolan-3-yl)-3-trifluoromethylphenyl]-1,3-oxazolan-5-ylmethyl} propanamide or its salts;

N1-((5S)-2-oxo-3-[4-(2-oxo-1,3-oxazolan-3-yl)-3-trifluoromethylphenyl]-1,3-oxazolan-5-ylmethyl} heptanamide or its salts;

N1-((5S)-2-oxo-3-[4-(2-oxo-1,3-oxazolan-3-yl)-3-

trifluoromethylphenyl]-1,3-oxazolan-5-ylmethyl} acrylamide or its salts;

N1-{(5S)-3-[2-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} acetamide or its salts;

N1-{(5S)-3-[3,5-difluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} acetamide or its salts;

N1-{(5S)-3-[3,5-difluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} propanamide or its salts;

N1-{(5S)-2-oxo-3-[4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-1,3-oxazolan-5-ylmethyl} acetamide or its salts;

N1-{(5S)-2-oxo-3-[4-(2-oxo-2,3-dihydrobenzo[d][1,3]oxazol-3-yl)phenyl]-1,3-oxazolan-5-ylmethyl} acetamide or its salts;

(5S)-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-5-(1-thioxoethylaminomethyl)-1,3-oxazolan-2-one or its salts;

(5S)-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-5-(3,3,3-trifluoro-1-thioxopropyl aminomethyl)-1,3-oxazolan-2-one or its salts;

3-{2-fluoro-4-[(5S)-2-oxo-5-(1-thioxoethylaminomethyl)-1,3-oxazolan-3-yl]phenyl}-2,3-dihydrobenzo[d][1,3]oxazol-2-one or its salts;

(5S)-3-[4-(2-oxo-1,3-oxazolan-3-yl)-3-trifluoromethylphenyl]-5-(1-thioxoethylamino methyl)-1,3-oxazolan-2-one or its salts;

(5S)-3-[3,5-difluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-5-(1-thioxoethylamino methyl)-1,3-oxazolan-2-one or its salts;

(5S)-3-[3,5-difluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-5-(1-thioxopropylamino methyl)-1,3-oxazolan-2-one or its salts;

(5S)-3-[4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-5-(1-thioxoethylaminomethyl)-1,3-oxazolan-2-one or its salts;

N1-{(5S)-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} methylcarbamate or its salts;

N1-{(5S)-2-oxo-3-[4-(2-oxo-1,3-oxazolan-3-yl)-3-trifluoromethylphenyl]-1,3-oxazolan-5-ylmethyl} methylcarbamate or its salts;

N1-{(5S)-3-[3,5-difluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} methylcarbamate or its salts;

N1-{(5S)-3-[4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} methyl carbamate or its salts;

(5S)-5-methylthioxy (thioxo)methylaminomethyl-3-[4-(2-oxo-1,3-oxazolan-3-yl) phenyl]-1,3-oxazolan-2-one or its salts;

N1-((5S)-3-[3,5-difluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} methylthiocarbamate or its salts;

N1-((5S)-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} methylthiocarbamate or its salts;

N1-((5S)-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} ethylthiocarbamate or its salts;

N1-((5S)-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} trifluoroacetoxythiocarbamate or its salts;

(5S)-5-cyclohexyloxy (thioxo)methylaminomethyl-3-[4-(2-oxo-1,3-oxazolan-3-yl) phenyl]-1,3-oxazolan-2-one or its salts;

N1-((5S)-3-[3-fluoro-4-(2-thioxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} methylthiocarbamate or its salts;

N1-((5S)-2-oxo-3-[4-(2-oxo-1,3-oxazolan-3-yl)-3-trifluoromethylphenyl]-1,3-oxazolan-5-ylmethyl} methylthiocarbamate or its salts;

N1-((5S)-2-oxo-3-[4-(2-oxo-1,3-oxazolan-3-yl)-3-trifluoromethylphenyl]-1,3-oxazolan-5-ylmethyl} ethylthiocarbamate or its salts;

N1-((5S)-3-[3,5-difluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} methylthiocarbamate or its salts;

N1-((5S)-3-[4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} methyl thiocarbamate or its salts;

N1-((5S)-3-[4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl} ethyl thiocarbamate or its salts;

N1-((5S)-3-[4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl}-1-propyl thiocarbamate or its salts;

(5S)-5-[2,2,2-trifluoroethyloxy(thioxo)methylaminomethyl]-3-[4-(2-oxo-1,3-oxazolan-3-yl) phenyl]-1,3-oxazolan-2-one or its salts;

(5S)-5-[2-hydroxyethyloxy (thioxo)methylaminomethyl]-3-[4-(2-oxo-1,3-oxazolan-3-yl) phenyl]-1,3-oxazolan-2-one or its salts;

(5S)-5-[2-methoxyethyloxy (thioxo)methylaminomethyl]-3-[4-(2-oxo-1,3-oxazolan-3-yl) phenyl]-1,3-oxazolan-2-one or its salts;

N1-((5S)-3-[4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-

oxazolan-5-ylmethyl} allylthio carbamate or its salts;

N1- {(5S)-3-[4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-2-oxo-1,3-oxazolan-5-ylmethyl}-2-propylthio carbamate or its salts;

N1- {(5S)-2-oxo-3-[4-(2-oxo-2,3-dihydrobenzo[d][1,3]oxazol-3-yl)phenyl]-1,3-oxazolan-5-ylmethyl} methylthiocarbamate or its salts;

N1- {(5S)-2-oxo-3-[4-(2-oxo-2,3-dihydrobenzo[d][1,3]oxazol-3-yl)phenyl]-1,3-oxazolan-5-ylmethyl} ethylthiocarbamate or its salts;

N1-((5S)-3-{4-[2-oxo-1,3-oxazolan-3yl]phenyl}-2-oxo-1,3-oxazolan-5-ylmethyl)-(N,N-dimethylamino)ethylthiocarbamate or its salts;

(5S)-5-[(2S)-2-hydroxymethylazolan-1-yl(thioxo)methylaminomethyl]-3-[4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-1,3-oxazolan-2-one or its salts;

(5S)-5-diethylamino (thioxo)methylaminomethyl-3-[4-(2-oxo-1,3-oxazolan-3-yl) phenyl]-1,3-oxazolan-2-one or its salts;

(5S)-5-allylamino (thioxo)methylaminomethyl-3-[4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-1,3-oxazolan-2-one or its salts;

(5S)-5-benzylamino(thioxo)methylaminomethyl-3-[4-(2-oxo-1,3-oxazolan-3-yl) phenyl]-1,3-oxazolan-2-one or its salts;

(5S)-5-[4-methoxybenzylamino(thioxo)methylaminomethyl]-3-[4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-1,3-oxazolan-2-one or its salts;

(5S)-3-[4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-5-[2-pyridylmethylamino (thioxo)methyl aminomethyl]-1,3-oxazolan-2-one or its salts;

(5S)-5-methylamino (thioxo)methylaminomethyl-3-[4-(2-oxo-1,3-oxazolan-3-yl) phenyl]-1,3-oxazolan-2-one or its salts;

(5S)-5-[2-hydroxyethylamino (thioxo)methylaminomethyl]-3-[4-(2-oxo-1,3-oxazolan-3-yl) phenyl]-1,3-oxazolan-2-one or its salts;

(5S)- 3-[4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-5-[1,4-thiazinan-4-yl(thioxo)methylamino methyl]-1,3-oxazolan-2-one or its salts;

(5S)-3-[4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-5-[2-pyridylamino (thioxo)methylamino methyl]- 1,3-oxazolan-2-one or its salts;

(5S)-5-amino(thioxo)methylaminomethyl-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-1,3-oxazolan-2-one or its salts;

(5S)-3-[3-fluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-5-methylamino (thioxo)methylamino methyl-1,3-oxazolan-2-one or its salts;

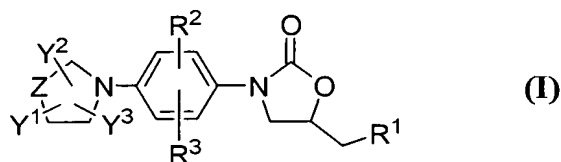
(5S)-5-amino(thioxo)methylaminomethyl-3-[3,5-difluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-1,3-oxazolan-2-one or its salts;

(5S)-3-[3,5-difluoro-4-(2-oxo-1,3-oxazolan-3-yl)phenyl]-5-methylamino(thioxo)methyl aminomethyl-1,3-oxazolan-2-one or its salts and

N1-((5S)-3-{4-[2-oxo-1,3-oxazolan-3yl]phenyl}-2-oxo-1,3-oxazolan-5-ylmethyl)-(N,N-dimethylamino)ethylthiocarbamate hydrochloride.

7. (Previously Presented) A compound according to claim 1, wherein the pharmaceutically acceptable salt is selected from Li, Na, K, Ca, Mg, Fe, Cu, Zn, or Mn or salts of organic bases, chiral bases, natural amino acids, unnatural amino acids, substituted amino acids, guanidine, substituted guanidine salts; ammonium, substituted ammonium salts, aluminum salts and acid addition salts.

8. (Withdrawn) A process for the preparation of the compound of formula (I)



where

R^1 represents NHR^4 , wherein R^4 represents hydrogen atom;

R^2 and R^3 are same or different and independently represent hydrogen, halogen atom, (C_1-C_6) alkyl group, halo (C_1-C_6) alkyl, cyano, nitro, SR^a , NR^a , OR^a where R^a represents substituted or unsubstituted (C_1-C_6) alkyl group, or halo (C_1-C_6) alkyl;

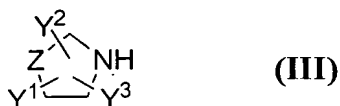
Z represents S or O;

Y^1 represents =O or =S group and Y^2 and Y^3 independently represent hydrogen, halogen, cyano, nitro, formyl, hydroxy, amino, =O, =S group, or substituted or unsubstituted groups selected from (C_1-C_6) alkyl, hydroxy (C_1-C_6) alkyl, (C_1-C_6) alkoxy (C_1-C_6) alkyl, (C_1-C_6) alkoxycarbonyl, carboxy (C_1-C_6) alkyl, (C_1-C_6) alkylsulfonyl, (C_1-C_6) alkylcarbonylamino (C_1-C_6) alkyl, arylcarbonylamino (C_1-C_6) alkyl, (C_1-C_6) alkylcarbonyloxy (C_1-C_6) alkyl, amino (C_1-C_6) alkyl, mono (C_1-C_6) alkylamino, di (C_1-C_6)

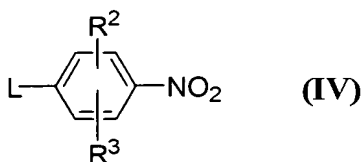
alkylamino, arylamino, (C₁-C₆)alkoxy, aryl, aryloxy, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocycloalkyl;

Y² and Y³ when present on adjacent carbon atoms together also form a substituted or unsubstituted 5 or 6 membered aromatic or non-aromatic cyclic structure, optionally containing one or two hetero atoms; its stereoisomers, its polymorphs, its pharmaceutically acceptable salts or its pharmaceutically acceptable solvates; which comprises:

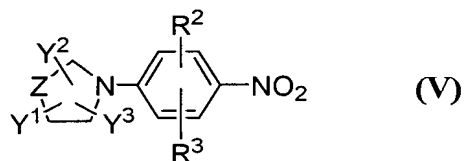
- (i) reacting a compound of formula (III)



where Y¹, Y², Y³ and Z are as defined above, with a compound of formula (IV)

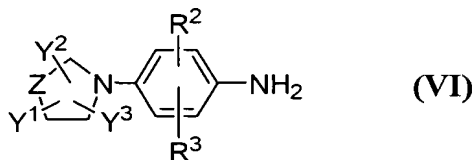


where L represents a leaving group; R² and R³ are as defined above, to produce a compound of formula (V)



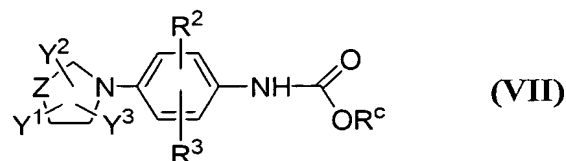
where Y¹, Y², Y³, R², R³ and Z are as defined above,

- (ii) reducing the compound of formula (V) to produce a compound of formula (VI)



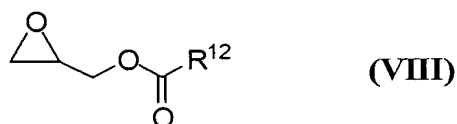
where Y¹, Y², Y³, R², R³ and Z are as defined above,

(iii) reacting the compound of formula (VI) with alkylchloroformate, to produce a compound of formula (VII)

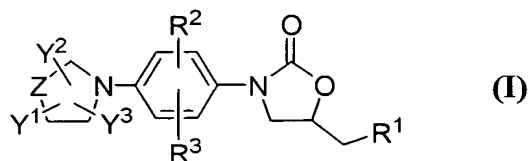


where R^c represents (C₁-C₈)alkyl group; Y¹, Y², Y³, R², R³ and Z are as defined above,

(iv) reacting the compound of formula (VII) with a compound of formula (VIII)

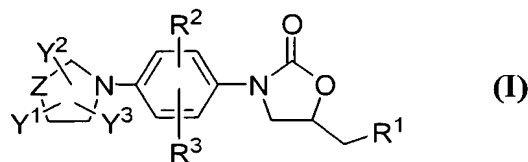


where R¹² represents (C₁-C₃)alkyl group in the presence of a base to produce a compound of formula (I)



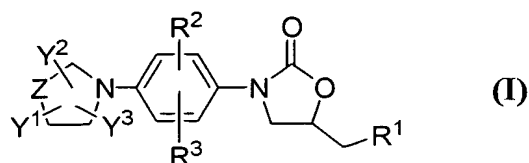
where R¹ represents hydroxy; Y¹, Y², Y³, R², R³ and Z are as defined above,

(v) reacting the compound of formula (I) with alkylsulfonyl chloride or aryl sulfonyl chloride to produce a compound of formula (I), where R¹ represents alkyl sulfonyl or aryl sulfonyl, and reacting with NaN₃ to produce compound of formula (I)



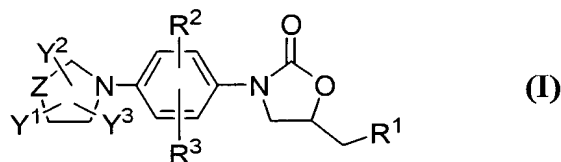
where R¹ represents azido group; Y¹, Y², Y³, R², R³ and Z are as defined above and

(vi) reducing the compound of formula (I) where R¹ represents azido group, to produce a compound of formula (I)



where R^1 represents NHR^4 wherein R^4 represents hydrogen atom; Y^1 , Y^2 , Y^3 , R^2 , R^3 and Z are as defined above.

9. (Withdrawn) A process for the preparation of compound of formula (I)



where

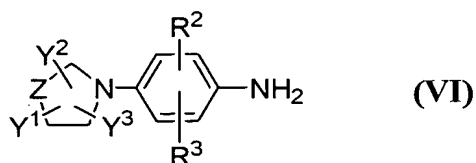
R^1 represents hydroxy;

R^2 and R^3 are same or different and independently represent hydrogen, halogen atom, (C_1-C_6) alkyl group, halo (C_1-C_6) alkyl, cyano, nitro, SR^a , NR^a , OR^a where R^a represents substituted or unsubstituted (C_1-C_6) alkyl group, or halo (C_1-C_6) alkyl; Z represents S or O;

Y^1 represents $=O$ or $=S$ group and Y^2 and Y^3 independently represent hydrogen, halogen, cyano, nitro, formyl, hydroxy, amino, $=O$, $=S$ group, or substituted or unsubstituted groups selected from (C_1-C_6) alkyl, hydroxy (C_1-C_6) alkyl, (C_1-C_6) alkoxy (C_1-C_6) alkyl, (C_1-C_6) alkoxycarbonyl, carboxy (C_1-C_6) alkyl, (C_1-C_6) alkylsulfonyl, (C_1-C_6) alkylcarbonylamino (C_1-C_6) alkyl, arylcarbonylamino (C_1-C_6) alkyl, (C_1-C_6) alkylcarbonyloxy (C_1-C_6) alkyl, amino (C_1-C_6) alkyl, mono (C_1-C_6) alkylamino, di (C_1-C_6) alkylamino, arylamino, (C_1-C_6) alkoxy, aryl, aryloxy, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocycloalkyl;

Y^2 and Y^3 when present on adjacent carbon atoms together may also form a substituted or unsubstituted 5 or 6 membered aromatic or non-aromatic cyclic structure, optionally containing one or two hetero atoms; its stereoisomers, its polymorphs, its pharmaceutically acceptable salts or its pharmaceutically acceptable solvates; which comprises:

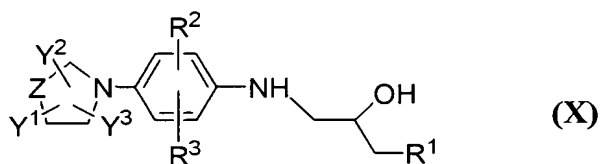
(i) reacting the compound of formula (VI)



where Y^1 , Y^2 , Y^3 , R^2 , R^3 and Z are as defined above, with a compound of formula (IX)



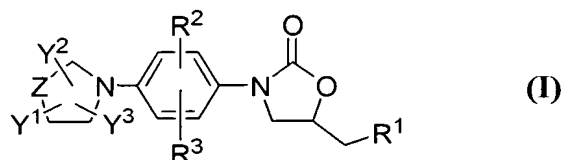
where R^1 represents hydroxy, to produce a compound of formula (X)



where R^1 represents hydroxy; Y^1 , Y^2 , Y^3 , R^2 , R^3 and Z are as defined above, and

(ii) carbonylating the compound of formula (X) with a carbonylating agent to produce the compound of formula (I) where R^1 represents hydroxy; Y^1 , Y^2 , Y^3 , R^2 , R^3 and Z are as defined above.

10. (Withdrawn) A process for the preparation of compound of the formula (I)



where

R^1 represents azido;

R^2 and R^3 are same or different and independently represent hydrogen, halogen atom, (C_1-C_6) alkyl group, halo (C_1-C_6) alkyl, cyano, nitro, SR^a , NR^a , OR^a where R^a represents substituted or unsubstituted (C_1-C_6) alkyl group, or halo (C_1-C_6) alkyl;

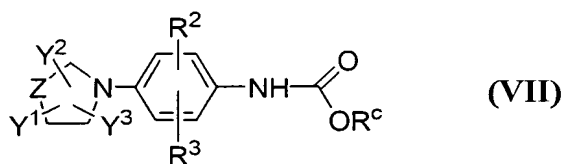
Z represents O;

Y^1 represents =O or =S group and Y^2 and Y^3 independently represent hydrogen, halogen, cyano, nitro, formyl, hydroxy, amino, =O, =S group, or substituted or unsubstituted groups selected from (C_1-C_6) alkyl, hydroxy (C_1-C_6) alkyl, (C_1-C_6)

alkoxy(C₁-C₆) alkyl, (C₁-C₆)alkoxycarbonyl, carboxy(C₁-C₆)alkyl, (C₁-C₆)alkylsulfonyl, (C₁-C₆)alkylcarbonylamino(C₁-C₆) alkyl, arylcarbonylamino(C₁-C₆)alkyl, (C₁-C₆)alkylcarbonyloxy(C₁-C₆)alkyl, amino(C₁-C₆)alkyl, mono(C₁-C₆)alkylamino, di(C₁-C₆)alkylamino, arylamino, (C₁-C₆) alkoxy, aryl, aryloxy, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocycloalkyl;

Y² and Y³ when present on adjacent carbon atoms together also form a substituted or unsubstituted 5 or 6 membered aromatic or non-aromatic cyclic structure, optionally containing one or two hetero atoms; its stereoisomers, its polymorphs, its pharmaceutically acceptable salts or its pharmaceutically acceptable solvates; which comprises:

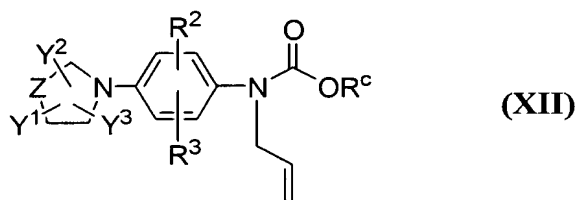
- (i) reacting a compound of formula (VII)



where R^c represents (C₁-C₈)alkyl group; Y¹, Y², Y³, R², R³ and Z are as defined above, with a compound of formula (XI)

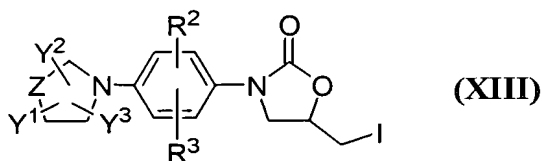


where L represents a leaving group; to produce a compound of formula (XII)



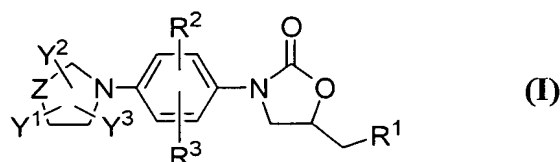
where R^c, Y¹, Y², Y³, R², R³ and Z are as defined above,

- (ii) converting the compound of formula (XII) defined above to a compound of formula (XIII)



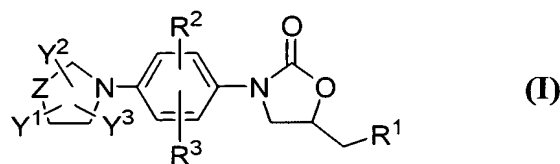
where Y^1 , Y^2 , Y^3 , R^2 , R^3 and Z are as defined above, and

(iii) converting the compound of formula (XIII) defined above to a compound of formula (I) by reacting with organic or inorganic azide



where R^1 represents azido group; Y^1 , Y^2 , Y^3 , R^2 , R^3 and Z are as defined above.

11. (Withdrawn) A process for the preparation of compound of formula (I)



where

R^1 represents azido group;

R^2 and R^3 are same or different and independently represent hydrogen, halogen atom, (C_1-C_6) alkyl group, halo (C_1-C_6) alkyl, cyano, nitro, SR^a , NR^a , OR^a where R^a represents substituted or unsubstituted (C_1-C_6) alkyl group, or halo (C_1-C_6) alkyl;

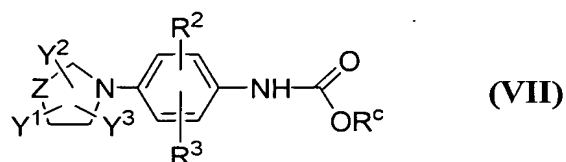
Z represents O;

Y^1 represents =O or =S group and Y^2 and Y^3 independently represent hydrogen, halogen, cyano, nitro, formyl, hydroxy, amino, =O, =S group, or substituted or unsubstituted groups selected from (C_1-C_6) alkyl, hydroxy (C_1-C_6) alkyl, (C_1-C_6) alkoxy (C_1-C_6) alkyl, (C_1-C_6) alkoxycarbonyl, carboxy (C_1-C_6) alkyl, (C_1-C_6) alkylsulfonyl, (C_1-C_6) alkylcarbonylamino (C_1-C_6) alkyl, arylcarbonylamino (C_1-C_6) alkyl, (C_1-C_6) alkylcarbonyloxy (C_1-C_6) alkyl, amino (C_1-C_6) alkyl, mono (C_1-C_6) alkylamino, di (C_1-C_6) alkylamino, arylamino, (C_1-C_6) alkoxy, aryl, aryloxy, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocycloalkyl;

Y^2 and Y^3 when present on adjacent carbon atoms together also form a substituted or unsubstituted 5 or 6 membered aromatic or non-aromatic cyclic structure, optionally containing one or two hetero atoms; its stereoisomers, its polymorphs,

its pharmaceutically acceptable salts or its pharmaceutically acceptable solvates; which comprises:

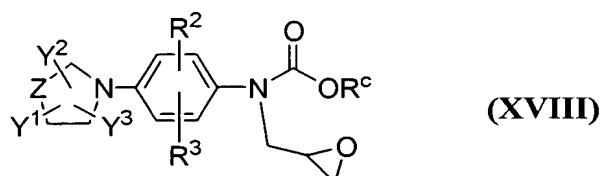
- (i) reacting a compound of formula (VII)



where R^c represents (C₁-C₈)alkyl group; Y¹, Y², Y³, R², R³ and Z are as defined above, with a compound of formula (XVII)

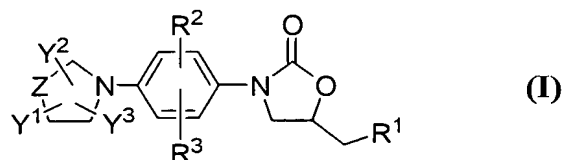


where L represents leaving group; to produce a compound of formula (XVIII)



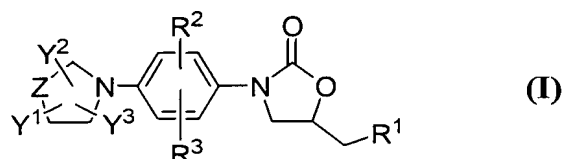
where R^c, Y¹, Y², Y³, R², R³ and Z are as defined above, and

- (ii) converting the compound of formula (XVIII) defined above to a compound of formula (I), by reacting with an organic or an inorganic azide,



where R¹ represents azido group; Y¹, Y², Y³, R², R³ and Z are as defined above.

12. (Withdrawn) A process for the preparation of compound of formula (I)



where

R^1 represents hydroxy group;

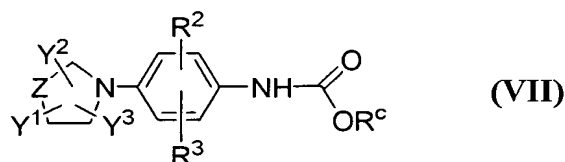
R^2 and R^3 are same or different and independently represent hydrogen, halogen atom, (C_1-C_6) alkyl group, halo (C_1-C_6) alkyl, cyano, nitro, SR^a , NR^a , OR^a where R^a represents substituted or unsubstituted (C_1-C_6) alkyl group, or halo (C_1-C_6) alkyl;

Z represents S or O;

Y^1 represents =O or =S group and Y^2 and Y^3 independently represent hydrogen, halogen, cyano, nitro, formyl, hydroxy, amino, =O, =S group, or substituted or unsubstituted groups selected from (C_1-C_6) alkyl, hydroxy (C_1-C_6) alkyl, (C_1-C_6) alkoxy (C_1-C_6) alkyl, (C_1-C_6) alkoxycarbonyl, carboxy (C_1-C_6) alkyl, (C_1-C_6) alkylsulfonyl, (C_1-C_6) alkylcarbonylamino (C_1-C_6) alkyl, arylcarbonylamino (C_1-C_6) alkyl, (C_1-C_6) alkylcarbonyloxy (C_1-C_6) alkyl, amino (C_1-C_6) alkyl, mono (C_1-C_6) alkylamino, di (C_1-C_6) alkylamino, arylamino, (C_1-C_6) alkoxy, aryl, aryloxy, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocycloalkyl;

Y^2 and Y^3 when present on adjacent carbon atoms together also form a substituted or unsubstituted 5 or 6 membered aromatic or non-aromatic cyclic structure, optionally containing one or two hetero atoms; its stereoisomers, its polymorphs, its pharmaceutically acceptable salts or its pharmaceutically acceptable solvates; which comprises:

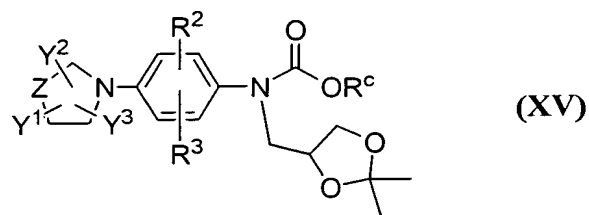
(i) reacting a compound of formula (VII)



where R^c represents (C_1-C_8) alkyl group; Y^1 , Y^2 , Y^3 , R^2 , R^3 and Z are as defined above, with a compound of formula (XIV)

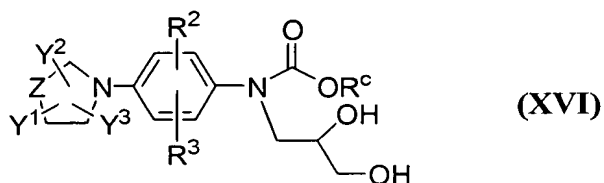


where L represents a leaving group; to produce a compound of formula (XV)



where R^c represents (C_1-C_8) alkyl group; Y^1 , Y^2 , Y^3 , R^2 , R^3 and Z are as defined above,

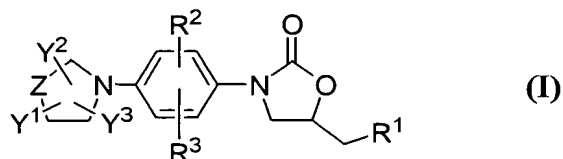
(ii) hydrolysing the acetonide moiety in the compound of formula (XV) to produce a compound of formula (XVI)



where R^c , Y^1 , Y^2 , Y^3 , R^2 , R^3 and Z are as defined above, and

(iii) cyclising the compound of formula (XVI) with or without a base to a compound of formula (I), where R^1 represents hydroxy group; Y^1 , Y^2 , Y^3 , R^2 , R^3 and Z are as defined above.

13. (Withdrawn) A process for the preparation of compound of the formula (I)



where

R^1 represents NHR^4 , wherein R^4 represents acetyl group;

R^2 and R^3 are same or different and independently represent hydrogen, halogen atom, (C_1-C_6) alkyl group, halo (C_1-C_6) alkyl, cyano, nitro, SR^a , NR^a , OR^a where R^a represents substituted or unsubstituted (C_1-C_6) alkyl group, or halo (C_1-C_6) alkyl;

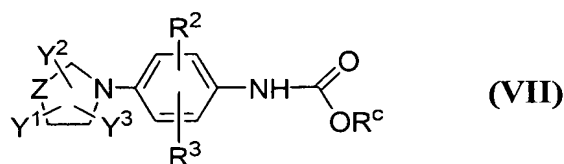
Z represents O;

Y^1 represents =O or =S group and Y^2 and Y^3 independently

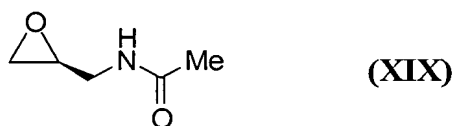
represent hydrogen, halogen, cyano, nitro, formyl, hydroxy, amino, =O, =S group, or substituted or unsubstituted groups selected from (C₁-C₆)alkyl, hydroxy(C₁-C₆)alkyl, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkoxycarbonyl, carboxy(C₁-C₆)alkyl, (C₁-C₆)alkylsulfonyl, (C₁-C₆)alkylcarbonylamino(C₁-C₆)alkyl, arylcarbonylamino(C₁-C₆)alkyl, (C₁-C₆)alkylcarbonyloxy(C₁-C₆)alkyl, amino(C₁-C₆)alkyl, mono(C₁-C₆)alkylamino, di(C₁-C₆)alkylamino, arylamino, (C₁-C₆)alkoxy, aryl, aryloxy, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocycloalkyl;

Y² and Y³ when present on adjacent carbon atoms together also form a substituted or unsubstituted 5 or 6 membered aromatic or non-aromatic cyclic structure, optionally containing one or two hetero atoms; its stereoisomers, its polymorphs, its pharmaceutically acceptable salts or its pharmaceutically acceptable solvates; which comprises:

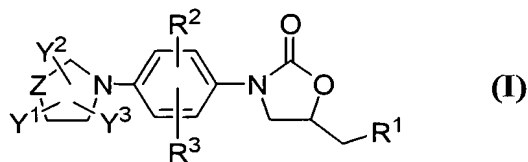
- (i) reacting a compound of formula (VII)



where R^c represents (C₁-C₈)alkyl group; Y¹, Y², Y³, R², R³ and Z are as defined above, with a compound of formula (XIX)

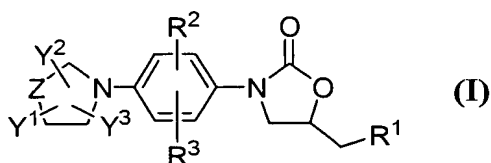


to produce a compound of formula (I)



where R¹ represents NHR⁴, wherein R⁴ represents acetyl group; Y¹, Y², Y³, R², R³ and Z are as defined above.

14. (Withdrawn) A process for the preparation of compound of formula (I)



where

R^1 represents NHR^4 , wherein R^4 represents formyl group;

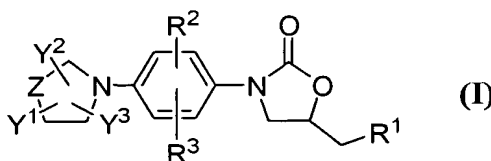
R^2 and R^3 are same or different and independently represent hydrogen, halogen atom, (C_1-C_6) alkyl group, halo (C_1-C_6) alkyl, cyano, nitro, SR^a , NR^a , OR^a where R^a represents substituted or unsubstituted (C_1-C_6) alkyl group, or halo (C_1-C_6) alkyl;

Z represents O;

Y^1 represents =O or =S group and Y^2 and Y^3 independently represent hydrogen, halogen, cyano, nitro, formyl, hydroxy, amino, =O, =S group, or substituted or unsubstituted groups selected from (C_1-C_6) alkyl, hydroxy (C_1-C_6) alkyl, (C_1-C_6) alkoxy (C_1-C_6) alkyl, (C_1-C_6) alkoxycarbonyl, carboxy (C_1-C_6) alkyl, (C_1-C_6) alkylsulfonyl, (C_1-C_6) alkylcarbonylamino (C_1-C_6) alkyl, arylcarbonylamino (C_1-C_6) alkyl, (C_1-C_6) alkylcarbonyloxy (C_1-C_6) alkyl, amino (C_1-C_6) alkyl, mono (C_1-C_6) alkylamino, di (C_1-C_6) alkylamino, arylamino, (C_1-C_6) alkoxy, aryl, aryloxy, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocycloalkyl;

Y^2 and Y^3 when present on adjacent carbon atoms together also form a substituted or unsubstituted 5 or 6 membered aromatic or non-aromatic cyclic structure, optionally containing one or two hetero atoms; which comprises: formylating the compound of formula (I) where R^1 represents NHR^4 wherein R^4 represents hydrogen and Y^1 , Y^2 , Y^3 , R^2 , R^3 and Z are as defined above, by using alkylformate.

15. (Withdrawn) A process for the preparation of compound of formula (I)

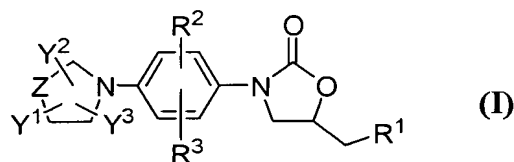


where

Y^1 represents =O or =S group and Y^2 and Y^3 independently represent hydrogen, halogen, cyano, nitro, formyl, hydroxy, amino, =O, =S group, or substituted or unsubstituted groups selected from (C₁-C₆)alkyl, hydroxy(C₁-C₆)alkyl, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkoxycarbonyl, carboxy(C₁-C₆)alkyl, (C₁-C₆)alkylsulfonyl, (C₁-C₆)alkylcarbonylamino(C₁-C₆)alkyl, arylcarbonylamino(C₁-C₆)alkyl, (C₁-C₆)alkylcarbonyloxy(C₁-C₆)alkyl, amino(C₁-C₆)alkyl, mono(C₁-C₆)alkylamino, di(C₁-C₆)alkylamino, arylamino, (C₁-C₆)alkoxy, aryl, aryloxy, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocycloalkyl;

Y^2 and Y^3 when present on adjacent carbon atoms together also form a substituted or unsubstituted 5 or 6 membered aromatic or non-aromatic cyclic structure, optionally containing one or two hetero atoms; its stereoisomers, its polymorphs, its pharmaceutically acceptable salts or its pharmaceutically acceptable solvates; which comprises: reacting compound of formula (I) where R^1 represents azido group; Y^1 , Y^2 , Y^3 , R^2 , R^3 and Z are as defined above, with thioacetic acid.

17. (Withdrawn) A process for the preparation of compound of formula (I)



where

R^1 represents NHR^4 , wherein R^4 represents $-C(=S)-R^{4b}$, wherein R^{4b} represents (C₁-C₆)alkyl, halo(C₁-C₆)alkyl, $-C(=O)-(C_1-C_6)$ alkoxy, $-C(=O)$ -aryloxy, $-C(=S)-(C_1-C_6)$ alkyl or $-C(=S)$ -aryl;

R^2 and R^3 are same or different and independently represent hydrogen, halogen atom, (C₁-C₆)alkyl group, halo(C₁-C₆)alkyl, cyano, nitro, SR^a , NR^a , OR^a where R^a represents substituted or unsubstituted (C₁-C₆)alkyl group, or halo(C₁-C₆)alkyl;

Z represents O;

Y^1 represents =O or =S group and Y^2 and Y^3 independently represent hydrogen, halogen, cyano, nitro, formyl, hydroxy, amino, =O, =S group, or substituted or unsubstituted groups selected from (C₁-C₆)alkyl, hydroxy(C₁-C₆)alkyl, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkoxycarbonyl, carboxy(C₁-C₆)alkyl, (C₁-C₆)alkylsulfonyl, (C₁-

Y² and Y³ when present on adjacent carbon atoms together also form a substituted or unsubstituted 5 or 6 membered aromatic or non-aromatic cyclic structure, optionally containing one or two hetero atoms; its stereoisomers, its polymorphs, its pharmaceutically acceptable salts or its pharmaceutically acceptable solvates; which comprises: reacting the compound of formula (I) where R¹ represents NHR⁴, wherein R⁴ represents -C(=O)-R^{4b}, wherein R^{4b} represents (C₁-C₆)alkyl, halo(C₁-C₆)alkyl, -C(=O)-(C₁-C₆)alkoxy, -C(=O)-aryloxy, -C(=S)-(C₁-C₆) alkyl or -C(=S)-aryl; Y¹, Y², Y³, R², R³ and Z are as defined above, with 2,4-bis (methoxyphenyl)-1,3-dithia-2,4-diphosphetane-2,4-disulfide (Lawesson's reagent).

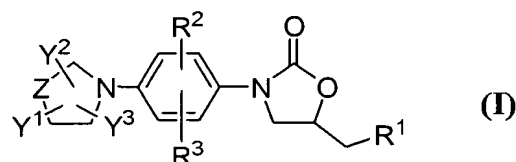
$$\text{Chemical structure (I): A central benzene ring substituted with a 1,3,4-oxadiazole ring (Y1, Y2, Y3, Z) and a 2-oxo-1,3-dihydroisoxazolidine ring (R1, R2, R3).}$$

Y¹ represents =O or =S group and Y² and Y³ independently represent hydrogen, halogen, cyano, nitro, formyl, hydroxy, amino, =O, =S group, or substituted or unsubstituted groups selected from (C₁-C₆)alkyl, hydroxy(C₁-C₆)alkyl, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkoxycarbonyl, carboxy(C₁-C₆)alkyl, (C₁-C₆)alkylsulfonyl, (C₁-C₆)alkylcarbonylamino(C₁-C₆)alkyl, arylcarbonylamino(C₁-C₆)alkyl, (C₁-C₆)alkylcarbonyloxy(C₁-C₆)alkyl, amino(C₁-C₆)alkyl, mono(C₁-C₆)alkylamino, di(C₁-C₆)

alkylamino, arylamino, (C₁-C₆) alkoxy, aryl, aryloxy, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocycloalkyl;

Y² and Y³ when present on adjacent carbon atoms together also form a substituted or unsubstituted 5 or 6 membered aromatic or non-aromatic cyclic structure, optionally containing one or two hetero atoms; its stereoisomers, its polymorphs, its pharmaceutically acceptable salts or its pharmaceutically acceptable solvates; which comprises: reacting the compound of formula (I) where R¹ represents NHR⁴, wherein R⁴ represents hydrogen; Y¹, Y², Y³, R², R³ and Z are as defined above, by reacting with carbondisulfide, with an alkylhalide and a base selected from Et₃N, diisopropylethylamine, K₂CO₃, NaH or KOt-Bu.

19. (Withdrawn) A process for the preparation of compound of formula (I)



where

R¹ represents NHR⁴, wherein R⁴ represents -C(=S)-OR^{4d}, wherein R^{4d} represents (C₁-C₆)alkyl, cyclo(C₃-C₆)alkyl, -(C=O)-(C₁-C₆)alkyl group substituted with fluorine; aryl, halo(C₁-C₆)alkyl, hydroxy(C₁-C₆)alkyl, (C₁-C₆)alkoxy(C₁-C₆)alkyl or (C₂-C₆)alkenyl;

R² and R³ are same or different and independently represent hydrogen, halogen atom, (C₁-C₆)alkyl group, halo(C₁-C₆)alkyl, cyano, nitro, SR^a, NR^a, OR^a where R^a represents substituted or unsubstituted (C₁-C₆)alkyl group, or halo(C₁-C₆)alkyl;

Z represents O;

Y¹ represents =O or =S group and Y² and Y³ independently represent hydrogen, halogen, cyano, nitro, formyl, hydroxy, amino, =O, =S group, or substituted or unsubstituted groups selected from (C₁-C₆)alkyl, hydroxy(C₁-C₆)alkyl, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkoxycarbonyl, carboxy(C₁-C₆)alkyl, (C₁-C₆)alkylsulfonyl, (C₁-C₆)alkylcarbonylamino(C₁-C₆)alkyl, arylcarbonylamino(C₁-C₆)alkyl, (C₁-C₆)alkylcarbonyloxy(C₁-C₆)alkyl, amino(C₁-C₆)alkyl, mono(C₁-C₆)alkylamino, di(C₁-C₆)

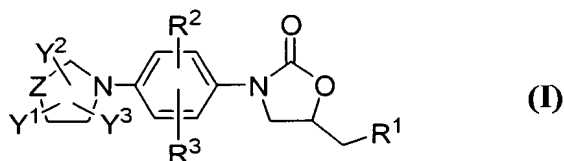
alkylamino, arylamino, (C₁-C₆) alkoxy, aryl, aryloxy, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocycloalkyl;

Y² and Y³ when present on adjacent carbon atoms together also form a substituted or unsubstituted 5 or 6 membered aromatic or non-aromatic cyclic structure, optionally containing one or two hetero atoms; its stereoisomers, its polymorphs, its pharmaceutically acceptable salts or its pharmaceutically acceptable solvates; which comprises:

(i) converting compound of formula (I) where R¹ represents NHR⁴, wherein R⁴ represents hydrogen atom; Y¹, Y², Y³, R², R³ and Z are as defined above, to a compound of formula (I) where R¹ represents isothiocyanate group and all other symbols are as defined above, by reacting with thiophosgene and

(ii) converting compound of formula (I) where R¹ represents isothiocyanate group, to a compound of formula (I) where R¹ represents NHR⁴, wherein R⁴ represents -C(=S)-OR^{4d}, wherein R^{4d} represents (C₁-C₆)alkyl, cyclo(C₃-C₆)alkyl, -(C=O)-(C₁-C₆) alkyl group substituted with fluorine; aryl, halo(C₁-C₆)alkyl, hydroxy(C₁-C₆)alkyl, (C₁-C₆) alkoxy(C₁-C₆)alkyl or (C₂-C₆)alkenyl and all symbols are as defined above, by reacting with alcohol.

20. (Withdrawn) A process for the preparation of compound of formula (I)



where

R¹ represents NHR⁴, wherein R⁴ represents -C(=S)-N(R'R''), R' represents hydrogen, (C₁-C₆)alkyl, (C₂-C₆)alkenyl, substituted or unsubstituted aralkyl, heteroaralkyl, hydroxy(C₁-C₆)alkyl, R'' represents hydrogen or (C₁-C₆)alkyl or R' and R'' together form a 5 or 6 membered cyclic structures containing one or two hetero atoms;

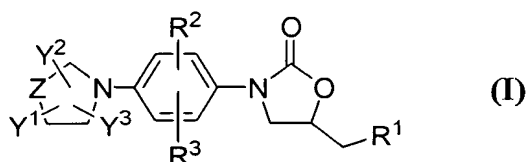
R² and R³ are same or different and independently represent hydrogen, halogen atom, (C₁-C₆)alkyl group, halo(C₁-C₆)alkyl, cyano, nitro, SR^a, NR^a, OR^a where R^a represents substituted or unsubstituted (C₁-C₆)alkyl group, or halo(C₁-C₆)alkyl; Z represents O;

Y^1 represents =O or =S group and Y^2 and Y^3 independently represent hydrogen, halogen, cyano, nitro, formyl, hydroxy, amino, =O, =S group, or substituted or unsubstituted groups selected from (C₁-C₆)alkyl, hydroxy(C₁-C₆)alkyl, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkoxycarbonyl, carboxy(C₁-C₆)alkyl, (C₁-C₆)alkylsulfonyl, (C₁-C₆)alkylcarbonylamino (C₁-C₆)alkyl, arylcarbonylamino(C₁-C₆)alkyl, (C₁-C₆)alkylcarbonyloxy(C₁-C₆)alkyl, amino(C₁-C₆)alkyl, mono(C₁-C₆)alkylamino, di(C₁-C₆)alkylamino, arylamino, (C₁-C₆)alkoxy, aryl, aryloxy, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocycloalkyl;

Y^2 and Y^3 when present on adjacent carbon atoms together also form a substituted or unsubstituted 5 or 6 membered aromatic or non-aromatic cyclic structure, optionally containing one or two hetero atoms; its stereoisomers, its polymorphs, its pharmaceutically acceptable salts or its pharmaceutically acceptable solvates; which comprises: converting compound of formula (I) where R¹ represents isothiocyanate group and all other symbols are as defined above by passing ammonia gas or by reacting with amine.

Claims 21 - 30. (Cancelled)

31. (Original) A pharmaceutical composition comprising a compound of formula (I)



as claimed in claim 1 and a pharmaceutically acceptable carrier, diluent, excipient or solvate.

32. (Original) A pharmaceutical composition as claimed in claim 31, in the form of a tablet, capsule, powder, syrup, solution or suspension.

33. (Previously presented) A method for treating a bacterial infection comprising administering an effective amount of a compound of formula (I) as claimed in claim 1 to a patient in need thereof.

34. (Previously Presented) A method for treating a bacterial infection comprising administering an effective amount of a pharmaceutical composition as claimed in claim 31 to a patient in need thereof.

35. (Original) A pharmaceutical composition comprising a compound as claimed in claim 6 and a pharmaceutically acceptable carrier, diluent, excipient or solvate.

36. (Original) A pharmaceutical composition as claimed in claim 35, in the form of a tablet, capsule, powder, syrup, solution or suspension.

37. (Previously presented) A method for treating a bacterial infection comprising administering an effective amount of a compound as claimed in claim 6 to a patient in need thereof.

38. (Previously presented) A method for treating a bacterial infection comprising administering an effective amount of a pharmaceutical composition as claimed in claim 35 to a patient in need thereof.

Claims 39 and 40. (Cancelled)

41. (Original) The compound according to claim 1, wherein the substituents on R^a are selected from hydroxy, halogen, nitro, amino, alkoxy, carboxy or cyano,

42. (Original) A compound according to claim 7, wherein the salts of organic bases are selected from N,N'-diacetylenediamine, betaine, caffeine, 2-diethylaminoethanol, 2-dimethylaminoethanol, N-ethylmorpholine, N-ethylpiperidine, glucamine, glucosamine, hydrabamine, isopropylamine, methylglucamine, morpholine, piperazine, piperidine, procaine, purines, theobromine, triethylamine, trimethylamine, tripropylamine, tromethamine, diethanolamine, meglumine, ethylenediamine, N,N'-diphenylethylenediamine, N,N'-dibenzylethylenediamine, N-benzyl phenylethylamine, choline, choline hydroxide, dicyclohexylamine, metformin, benzylamine, phenylethylamine, dialkylamine, trialkylamine, thiamine, aminopyrimidine, aminopyridine, purine, or spermidine.

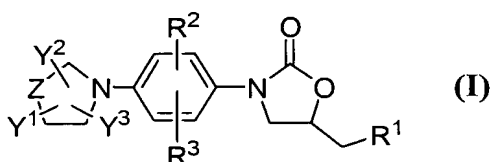
43. (Original) A compound according to claim 7, wherein the salts of chiral bases are selected from alkylphenylamine, glycinol, phenyl glycinol.

44. (Original) A compound according to claim 7, wherein the salts of natural amino acids are selected from glycine, alanine, valine, leucine, isoleucine, norleucine, tyrosine, cystine, cysteine, methionine, proline, hydroxy proline, histidine, ornithine, lysine, arginine, serine, threonine, or phenylalanine.

45. (Original) A compound according to claim 7, wherein the salts of unnatural amino acid, substituted amino acids are selected from D-isomers, guanidine, substituted guanidine wherein the substituents are selected from nitro, amino, alkyl selected from methyl, ethyl, and propyl; alkenyl selected from ethenyl, propenyl, or butenyl; alkynyl selected from ethynyl, or propynyl.

46. (Original) A compound according to claim 7, wherein the addition salts are selected from sulphates, nitrates, phosphates, perchlorates, borates, halides, acetates, tartrates, maleates, citrates, succinates, palmoates, methanesulphonates, benzoates, salicylates, hydroxynaphthoates, benzenesulfonates, ascorbates, glycerophosphates, or ketoglutarates.

47. (Withdrawn) A process for the preparation of compound of formula (I),



where

R^1 represents halogen atom;

R^2 and R^3 are same or different and independently represent hydrogen, halogen atom, (C_1-C_6) alkyl group, halo (C_1-C_6) alkyl, cyano, nitro, SR^a , NR^a , OR^a where R^a represents substituted or unsubstituted (C_1-C_6) alkyl group, or halo (C_1-C_6) alkyl;

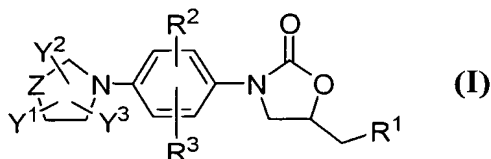
Z represents O;

Y^1 represents =O or =S group and Y^2 and Y^3 independently

represent hydrogen, halogen, cyano, nitro, formyl, hydroxy, amino, =O, =S group, or substituted or unsubstituted groups selected from (C₁-C₆)alkyl, hydroxy(C₁-C₆)alkyl, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkoxycarbonyl, carboxy(C₁-C₆)alkyl, (C₁-C₆)alkylsulfonyl, (C₁-C₆)alkylcarbonylamino(C₁-C₆)alkyl, arylcarbonylamino(C₁-C₆)alkyl, (C₁-C₆)alkylcarbonyloxy(C₁-C₆)alkyl, amino(C₁-C₆)alkyl, mono(C₁-C₆)alkylamino, di(C₁-C₆)alkylamino arylamino, (C₁-C₆)alkoxy, aryl, aryloxy, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocycloalkyl;

Y² and Y³ when present on adjacent carbon atoms together also form a substituted or unsubstituted 5 or 6 membered aromatic or non-aromatic cyclic structure, optionally containing one or two hetero atoms; its stereoisomers, its polymorphs, its pharmaceutically acceptable salts or its pharmaceutically acceptable solvates; which comprises: reacting a compound of formula (I) where R¹ represents hydroxy group and all other symbols are as defined above, with tetrahalomethane group and PPh₃ or P(alkyl)₃.

48. (Withdrawn) A process for the preparation of compound of formula (I),



where

R¹ represents SH group;

R² and R³ are same or different and independently represent hydrogen, halogen atom, (C₁-C₆)alkyl group, halo(C₁-C₆)alkyl, cyano, nitro, SR^a, NR^a, OR^a where R^a represents substituted or unsubstituted (C₁-C₆)alkyl group, or halo(C₁-C₆)alkyl;

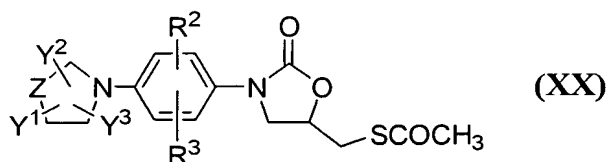
Z represents O;

Y¹ represents =O or =S group and Y² and Y³ independently represent hydrogen, halogen, cyano, nitro, formyl, hydroxy, amino, =O, =S group, or substituted or unsubstituted groups selected from (C₁-C₆)alkyl, hydroxy(C₁-C₆)alkyl, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkoxycarbonyl, carboxy(C₁-C₆)alkyl, (C₁-C₆)alkylsulfonyl, (C₁-C₆)alkylcarbonylamino(C₁-C₆)alkyl, arylcarbonylamino(C₁-C₆)alkyl, (C₁-C₆)alkylcarbonyloxy(C₁-C₆)alkyl, amino(C₁-C₆)alkyl, mono(C₁-C₆)alkylamino, di(C₁-C₆)

alkylamino arylamino, (C₁-C₆) alkoxy, aryl, aryloxy, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocycloalkyl;

Y² and Y³ when present on adjacent carbon atoms together also form a substituted or unsubstituted 5 or 6 membered aromatic or non-aromatic cyclic structure, optionally containing one or two hetero atoms; its stereoisomers, its polymorphs, its pharmaceutically acceptable salts or its pharmaceutically acceptable solvates; which comprises:

(i) reacting the compound of formula (I) where R¹ represents halogen atom, to produce a compound of formula (XX),



where all symbols are as defined above, with a base and thioacetic acid,

(ii) reacting the compound of formula (XX), to produce a compound of formula (I) where R¹ represents SH group and all other symbols are as defined earlier, with base.

49. (Previously presented) A compound according to claim 6, wherein the pharmaceutically acceptable salt is selected from Li, Na, K, Ca, Mg, Fe, Cu, Zn, or Mn or salts of organic bases, chiral bases, natural amino acids, unnatural amino acids, substituted amino acids, guanidine, substituted guanidine salts; ammonium, substituted ammonium salts, aluminum salts and acid addition salts.

50. (Previously presented) A method for treating a bacterial infection comprising administering an effective amount of a pharmaceutical composition as claimed in claim 32 to a patient in need thereof.

51. (Previously presented) A method for treating a bacterial infection comprising administering a pharmaceutical composition as claimed in claim 36 to a patient in need thereof.